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## Amendments to the Claims:

This listing of claims replaces all prior versions and listings of claims in the application:

## **Listing of Claims**:

1. (Currently amended) A method for analysis of proteins in a biological system comprising:

providing a biological system;

exposing the system to a stimulus;

sampling the biological system at multiple time intervals-to provide multiple samples, each sample containing multiple proteinsafter exposing the system to the stimulus;

treating submitting each of the multiple samples byto a separation technique to provide multiple protein samples suitable for analysis by mass spectrometry; and

analyzing the multiple samples to determine changes in protein abundance of proteins as a function of time after exposing the biological system to stimulus, said analyzing including

allocating the multiple protein samples for the multiple samples among mass spectrometry systems in a parallel array of mass spectrometry systems, each mass spectrometry system adapted for protein analysis and providing mass spectral data indicating identity and abundance of one or more proteins, providing a parallel array of mass spectrometry systems adapted for protein analysis, and

directing mass spectral data from <u>each of</u> the mass spectrometry systems in said array to a common computing device, said mass spectral data being indicative of the identity and the abundance of protein in said multiple sample, and

<u>correlating collating</u> said mass spectral data <u>from each of the mass spectrometry</u> <u>systems</u> as a function of time <u>of sampling of the biological system</u>.

2. (Currently amended) The method of claim 1 <u>further</u> comprising displaying said co<u>llated</u>rrelated data as a function of protein identity, protein abundance, and time.

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3. (Canceled)

4. (Canceled)

- 5. (Currently amended) The method of claim [[4]]1 wherein said array of mass spectrometry systems includes at least 20-5 mass spectrometers.
- 6. (Currently amended) The method of claim [[4]]1 wherein comprising analyzing the multiple samples includes analyzing the multiple samples to determine changes in abundance of 500 proteins or more.
- 7. (Currently amended) The method of claim [[6]]1 comprising wherein analyzing the multiple samples includes analyzing the multiple samples to determine changes in abundance of about 5000 proteins or more.
- 8. (Currently amended) The method of claim [[4]]1 wherein the separation technique includes use of one or more separation apparatus and said common computing device communicates with each of said separation apparatus.
- 9. (Currently amended) The method of claim [[8]]1 wherein the separation technique includes use of liquid chromatography.
- 10. (Currently amended) The method of claim 8 wherein the separation technique apparatus includes use of a magnetic particle separation apparatus.
- 11. (Currently amended) The method of claim [[10]]38 wherein the magnetic particlearray of separation apparatus treats multiple samples in parallel.

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12. (Currently amended) The method of claim [[4]]1 wherein the separation technique includes treating each of the multiple protein samples with a protease to produce peptides and the mass spectral data includes said mass spectral data includes peptide fragment mass spectra and an amino acid sequence data that can be compared to amino acid sequence data derived from a data base.

- 13. (Currently amended) The method of claim 12 wherein said mass spectrometry systemser are LC-TMS mass spectrometers.
- 14. (Currently amended) The method of claim [[4]]1 <u>further comprising:</u>
  exposing a first <u>eomponentinstance</u> of the biological system to a stimulus and maintaining a second <u>eomponent instance</u> of the biological system free of the stimulus; wherein

sampling, submitting, and analyzing include sampling, submitting, and analyzing each of the first component and the second instances component; and

<u>correlating mass spectral data includes comparing the identity and abundance mass</u>
<u>spectral data from in the first component and the second component instances.</u>

- 15. (Original) The method of claim 14 comprising separately analyzing samples from said first component and second component.
- 16. (Currently amended) The method of claim [[4]]43 wherein the stimulus perturbation results from exposure of the biological system to heat, light, cold, motion, agitation, cellular material, or a is a drug.
- 17. (Currently amended) The method of claim [[4]]1 wherein the time interval is about 5 to 60 seconds.

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18. (Currently amended) The method of claim [[4]]1 wherein the time interval is about one minute to one hour.

Claims 19-21. (Cancelled)

22. (Currently amended) A method for analysis of proteins in a biological system comprising:

providing a biological system containing proteins;

exposing the biological system to a stimulus;

after exposing the biological system to the stimulus, sampling the biological system at multiple time intervals to obtain multiple samples, each sample containing multiple proteins;

treating the multiple samples by a <u>parallel</u> separation technique to provide multiple protein samples suitable for analysis by mass spectrometry;

providing a parallel array of mass spectrometer systems capable of simultaneous analysis of as many protein samples as there are spectrometer systems in said array;

analyzing allocating the multiple protein samples among the mass spectrometry systems in the parallel array of mass spectrometry systems in said parallel array of mass spectrometry systems to generate obtain mass spectral data indicating of the identity and the abundance of proteins in said multiple protein samples; and

communicating the mass spectral data to in-a common electronic computing device communicating with each of said mass spectrometry systems, and

correlating said mass spectral data as a function of time.

23. (Currently amended) The method of claim 22 where<u>in</u> the parallel separation device <u>technique</u> is <u>performed using</u> a parallel magnetic particle separation device.

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24. (Currently amended) The method of claim 23 wherein the parallel array of mass spectrometry systems includes an array of LC-MS spectrometer systems.

- 25. (Previously presented) The method of claim 24 wherein the array includes 6-20 mass spectrometers.
- 26. (Previously presented) The method of claim 25 wherein the time intervals are in the range of 5 seconds to 10 minutes.
- 27. (Previously presented) The method of claim 26 wherein the analysis includes analysis of about 500 proteins or more.
- 28. (Currently amended) The method of claim 2<u>3</u>7 wherein the central computer communicates with the parallel magnetic particle separation device separation.
- 29. (Currently amended) A system for mass spectrometric analysis of proteins in a biological system, the system comprising:

a parallel sample separation apparatus adapted to receive multiple samples of a biological system taken at multiple time intervals, and separate the multiple samples in parallel to obtain multiple protein samples for analysis by mass spectrometry;

a parallel array of mass spectrometry systems adapted to receive the <u>multiple</u> protein samples from the separation apparatus and analyze the <u>received multiple</u> protein samples in parallel to generate mass spectral data indicatingive of the identity and the abundance of proteins in the multiple protein samples; and

a computing device communicating with the parallel array of mass spectrometry systems and the parallel separation apparatus, the computing device being adapted to analyze the mass spectral data from the parallel array of mass spectrometry systems and collaterrelate the mass spectral data as a function of time of sampling.

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30. (Previously presented) The system of claim 29, wherein the parallel separation device is a parallel magnetic particle separation device.

- 31. (Previously presented) The system of claim 29, wherein the parallel separation device is a parallel chromatography separation device.
- 32. (Currently amended) The system of claim 29, wherein the computing device is adapted to collaterrelate the mass spectral data as a function of protein identity, protein abundance, and time.
- 33. (Currently amended) The system of claim 29, further comprising a searchable database adapted to storegraphical user interface that can be searched, queried, or filtered to display selected the correlated collated data.
- 34. (Currently amended) The system of claim 29 wherein the parallel array of mass spectrometry systems includes at least 2-5 mass spectrometers.
- 35. (Previously presented) The system of claim 29, wherein the parallel array of mass spectrometry systems includes at least 20 mass spectrometers.
- 36. (Previously presented) The system of claim 29, wherein the parallel array of mass spectrometry systems is adapted to generate mass spectral data including peptide fragment mass spectra, and the computing device is adapted to analyze the mass spectral data in conjunction with an amino acid sequence derived from a database.
- 37. (Previously presented) The system of claim 29, wherein the parallel array of mass spectrometry systems include a liquid chromatograph-tandem mass spectrometer system.

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38. (New) The method of claim 8 wherein a first portion of the multiple protein samples are allocated among the mass spectrometry systems before a second portion of the multiple protein samples have been provided by the separation technique.

- 39. (New) The method of claim 8 wherein the separation technique includes use of an array of parallel separation apparatus.
- 40. (New) The method of claim 39 wherein the number of separation apparatus in the array of parallel separation apparatus is equal to the number of mass spectrometry systems in the array of parallel mass spectrometry systems.
- 41. (New) The method of claim 11 wherein the array of mass spectrometry systems treat multiple samples in parallel.
- 42. (New) The method of claim 41 wherein treatment of multiple samples by the array of separation apparatus is carried out in parallel with treatment of multiple samples by the array of mass spectrometry systems.
- 43. (New) The method of claim 1 further comprising exposing the biological system to a perturbation, wherein sampling of the biological system occurs at multiple time intervals after the exposure of the biological system to the perturbation.
- 44. (New) The method of claim 1 further comprising inferring interactions over time between and among proteins in the biological system.
- 45. (New) The method of claim 2 wherein protein abundance is expressed as relative abundance of proteins in each of the multiple samples.